Amendments to the Claims

This listing of claims will replace all prior versions and listings of claims in the application:

(currently amended) A method for treating allergic rhinitis in mammals which comprises intranasally administering a pharmaceutically effective amount of a composition comprising 0.01 0.1 - 0.8 % (w/v) olopatadine and 0.01 - 1.0 0.02 - 0.5 % (w/v) of a steroid selected from the group consisting of fluticasone, mometasone, budesonide and beclomethasone, wherein the composition has a pH of 3.5 - 8.0 and a viscosity of 1 - 50 cps.

2 & 3. (cancelled).

- 4. (previously presented) The method of Claim 1 wherein the steroid is fluticasone.
- 5. (original) The method of Claim 1 wherein the steroid has an average particle size of $2.5 5 \mu m$.
- 6. (original) The method of Claim 1 wherein the steroid has an average particle size of less than 0.8 μ m.
- 7. (original) The method of Claim 6 wherein the steroid has an average particle size of $0.5 \, \mu \text{m}$ or less.
- 8. (original) The method of Claim 1 wherein the composition is an aqueous composition packaged as a nasal spray.
- 9. (cancelled).
- 10. (currently amended) A method for treating allergic rhinitis in mammals which comprises intranasally administering a pharmaceutically effective amount of a composition comprising 0.01 0.4 0.8 % (w/v) olopatadine and 0.02 0.5 % (w/v) of a steroid selected from the group consisting of fluticasone, mometasone, budesonide and beclomethasone, wherein the composition has a pH of 3.5 8.0 and

U.S. Serial No. 10/706,759 Filed: November 12, 2003

a viscosity of 1-50 cps., and the composition is an aqueous composition packaged as a nasal spray.